

STN Search History

FILE 'HOME' ENTERED AT 08:43:37 ON 28 FEB 2006

- L1 2956 (VACCINE OR DTP OR DPT OR BOOSTER) AND (REDUC? OR LOW?) (S)
(ALUM? OR DIPHTHERIA OR TETAN? OR TOXIN# OR TOXOID#)
- L7 8 L6 AND (DIPHTHERIA OR TETAN? OR DTP OR DPT) AND (VACCINE OR
BOOSTER)

(FILE 'HOME' ENTERED AT 08:43:37 ON 28 FEB 2006)

FILE 'MEDLINE, CAPLUS, BIOSIS, EMBASE, SCISEARCH' ENTERED AT 08:44:02 ON
28 FEB 2006

- L1 2956 S (VACCINE OR DTP OR DPT OR BOOSTER) AND (REDUC? OR LOW?) (S) (
- L2 768 S L1 AND (DIPHTHERIA (P) TETAN? (P) (PERTUSSIS OR POLIO?))
- L3 1564 S L1 AND (REDUC? OR LOW?) (S) (DIPHTHERIA OR TETAN?)
- L4 212 S L1 AND (REDUC? OR LOW?) (5N) (ALUM?)
- L5 85 DUP REM L4 (127 DUPLICATES REMOVED)
- L6 57 S L5 AND PY<1999
- L7 8 S L6 AND (DIPHTHERIA OR TETAN? OR DTP OR DPT) AND (VACCINE OR
- L8 49 S L6 NOT L7
- L9 0 S L8 AND (LOW? OR REDUC?) (S) TETAN? (S) DIPHTH?
- L10 3 S L8 AND (DTP? OR DTAP OR ADTP OR DPT)
- L11 647 S L3 AND (LOW? OR REDUC?) (S) DIPHTH? (S) TETAN?
- L12 522 S L11 AND L2
- L13 232 S L12 AND PY<1999
- L14 258 S L3 AND (LOW? OR REDUC?) (6N) DIPHTH? (6N) TETAN?
- L15 75 S L13 AND L14
- L16 43 DUP REM L15 (32 DUPLICATES REMOVED)
- L17 417 S L3 AND (REDUC? OR LOW?) (5N) (DIPHTH? OR TETAN?) NOT L11
- L18 189 DUP REM L17 (228 DUPLICATES REMOVED)
- L19 120 S L18 AND PY<1999
- L20 15 S L19 AND ((REDUC? OR LOW?) (5N) (TETAN? OR DIPHTH?))/TI

L7 ANSWER 1 OF 8 MEDLINE on STN
 AN 97185150 MEDLINE
 DN PubMed ID: 9032891
 TI Single shot with **tetanus** toxoid in biodegradable microspheres protects mice despite acid-induced denaturation of the antigen.
 AU Kersten G F; Donders D; Akkermans A; Beuvery E C
 CS National Institute of Public Health and Environmental Protection, Department of Product and Process Development, Bilthoven, The Netherlands.
 SO Vaccine, (1996 Dec) Vol. 14, No. 17-18, pp. 1627-32.
 Journal code: 8406899. ISSN: 0264-410X.
 CY ENGLAND: United Kingdom
 DT Journal; Article; (JOURNAL ARTICLE)
 LA English
 FS Priority Journals
 EM 199705
 ED Entered STN: 19970523
 Last Updated on STN: 19970523
 Entered Medline: 19970509
 AB **Tetanus** toxoid encapsulated in microspheres consisting of biodegradable polyesters, prepared by four different manufacturers were evaluated with respect to antigenic load, in vitro release pattern, antigen integrity and immunogenicity. In vitro release studies over periods up to 140 days indicated that only during the first days **tetanus** toxoid was released. Although some preparations were designed to release their antigen content in a pulsatile manner, this was never observed in vitro. A single immunization with 0.3 Lf **tetanus** toxoid in microspheres induced substantial humoral responses, in most cases higher than one immunization with plain **tetanus** toxoid, sometimes higher than one dose of alum-adsorbed toxoid but always lower than booster immunizations. It is shown that the moderate (no booster effect) performance of the microsphere preparations is probably due to acid induced denaturation of the antigen. Despite this drawback, protection level in mice after challenge with 50 LD50 1 year after one immunization with microspheres was, on average, substantially higher than mice receiving plain **tetanus** toxoid.

L7 ANSWER 2 OF 8 MEDLINE on STN
 AN 96155147 MEDLINE
 DN PubMed ID: 8585294
 TI Adjuvant activity of aluminium hydroxide and calcium phosphate in **diphtheria-tetanus vaccines--I**.
 AU Aggerbeck H; Heron I
 CS Statens Serum Institut, Bacterial Vaccine Department, Copenhagen, Denmark.
 SO Vaccine, (1995 Oct) Vol. 13, No. 14, pp. 1360-5.
 Journal code: 8406899. ISSN: 0264-410X.
 CY ENGLAND: United Kingdom
 DT Journal; Article; (JOURNAL ARTICLE)
 LA English
 FS Priority Journals
 EM 199603
 ED Entered STN: 19960327
 Last Updated on STN: 19970203
 Entered Medline: 19960319
 AB The potencies of two **diphtheria-tetanus vaccines** (DT) adsorbed to either aluminium hydroxide or calcium phosphate were compared in mice and guinea pigs. The **vaccines** were made from the same batches of purified toxoids and contained the same amounts of antigens. Immunizations were done once or twice with different doses of **vaccine** injected undiluted, diluted in saline or diluted in the corresponding adjuvant. The various potency assays showed that the adjuvant activity of calcium phosphate was lower than or equal to aluminium hydroxide. Despite the range of potency assays done, none of the methods reflected the efficacy of these **vaccines** in revaccination of humans. A simplified potency assay is suggested for release of final **vaccine** formulations to reduce the number of animals in quality control.

L7 ANSWER 7 OF 8 BIOSIS COPYRIGHT (c) 2006 The Thomson Corporation on STN
AN 1995:275303 BIOSIS
DN PREV199598289603
TI Can **reductions** in **diphtheria toxoid** or
aluminum content **reduce** the reactogenicity of
booster doses of **DPT vaccine**?.
AU Scheifele, David W. [Reprint author]; Meekison, William; Guasparini,
Roland; Barreto, Luis
CS Vaccine Eval. Cent., 950 West 28th Ave., Vancouver, BC V5Z 4H4, Canada
SO Immunology and Infectious Diseases (Oxford), (1995) Vol. 5, No.
1, pp. 73-77.
CODEN: IINDEK. ISSN: 0959-4957.
DT Article
LA English
ED Entered STN: 26 Jun 1995
Last Updated on STN: 26 Jun 1995
AB Large local reactions often follow **booster** doses of the
diphtheria-pertussis-tetanus (DPT)
vaccine used in Canada. Their cause is uncertain but the
relatively large dose of **diphtheria toxoid** is a possible
contributor. This study was designed to determine if **vaccines**
containing less **diphtheria toxoid** (25, 15 or 10 Lf per dose)
and/or aluminum (0.33 or 0.17 mg per dose) would cause fewer adverse
effects. Subjects included 373 healthy 4- to 6-year-olds due for a fifth
dose of **DPT vaccine**. They were randomly assigned to
receive one of five **DPT vaccines**, in double-blinded
fashion. Local and systemic adverse reactions were assessed 24 h
post-immunization by visiting research nurses. Blood was tested for
antitoxins prior to and 4 weeks after immunization. The lot of standard
vaccine caused fever in 21.9% and gtoreq 50 mm of injection site
redness in 49.3% and swelling in 41.0%. Limited arm movement was detected
in 26% and was severe in 5.5%. None of the modified **vaccines**
reduced such morbidity significantly. All formulations produced large
increases in **tetanus** and **diphtheria** antitoxin levels.
It is concluded that the principal cause of local reactions after
vaccine is not **diphtheria toxoid** or aluminum but a
component(s) that was not varied among the five preparations, such as
pertussis **vaccine**. The advent of acellular pertussis
vaccines might provide a remedy.

L16 ANSWER 1 OF 43 CAPLUS COPYRIGHT 2006 ACS on STN

AN 1998:323156 CAPLUS

DN 129:19687

TI Acellular **pertussis vaccine** with **diphtheria**
and **tetanus** toxoids

IN Florent, Patrick; Stephenne, Jean; Vandecasserie, Christian

PA Smithkline Beecham Biologicals SA, Belg.; Florent, Patrick; Stephenne,
Jean; Vandecasserie, Christian

SO PCT Int. Appl., 26 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

| | PATENT NO. | KIND | DATE | APPLICATION NO. | DATE |
|------|---|------|----------|------------------|--------------|
| PI | WO 9819702 | A1 | 19980514 | WO 1997-EP6180 | 19971104 <-- |
| | W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW | | | | |
| | RW: GH, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG | | | | |
| | CA 2271008 | AA | 19980514 | CA 1997-2271008 | 19971104 <-- |
| | AU 9853196 | A1 | 19980529 | AU 1998-53196 | 19971104 <-- |
| | AU 710475 | B2 | 19990923 | | |
| | EP 941117 | A1 | 19990915 | EP 1997-950137 | 19971104 |
| | EP 941117 | B1 | 20020828 | | |
| | R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, FI | | | | |
| | CN 1236321 | A | 19991124 | CN 1997-199491 | 19971104 |
| | CN 1130226 | B | 20031210 | | |
| | BR 9712917 | A | 19991207 | BR 1997-12917 | 19971104 |
| | NZ 335384 | A | 20001027 | NZ 1997-335384 | 19971104 |
| | JP 2001503422 | T2 | 20010313 | JP 1998-521070 | 19971104 |
| | AT 222773 | E | 20020915 | AT 1997-950137 | 19971104 |
| | EP 1240905 | A1 | 20020918 | EP 2002-75821 | 19971104 |
| | EP 1240905 | B1 | 20050511 | | |
| | R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, FI | | | | |
| | PT 941117 | T | 20021231 | PT 1997-950137 | 19971104 |
| | ES 2182131 | T3 | 20030301 | ES 1997-950137 | 19971104 |
| | PL 188460 | B1 | 20050228 | PL 1997-333127 | 19971104 |
| | AT 295178 | E | 20050515 | AT 2002-75821 | 19971104 |
| | PT 1240905 | T | 20050930 | PT 2002-75821 | 19971104 |
| | ES 2240647 | T3 | 20051016 | ES 2002-75821 | 19971104 |
| | CZ 295954 | B6 | 20051214 | CZ 1999-1640 | 19971104 |
| | ZA 9709984 | A | 19980723 | ZA 1997-9984 | 19971106 <-- |
| | TW 471971 | B | 20020111 | TW 1997-86119712 | 19971224 |
| | NO 9902156 | A | 19990504 | NO 1999-2156 | 19990504 |
| | NO 320186 | B1 | 20051107 | | |
| | KR 2000053092 | A | 20000825 | KR 1999-704016 | 19990506 |
| | MX 9904278 | A | 20000131 | MX 1999-4278 | 19990507 |
| | HK 1022638 | A1 | 20030228 | HK 2000-101495 | 20000310 |
| | US 2001014331 | A1 | 20010816 | US 2001-827785 | 20010406 |
| | US 2004208898 | A1 | 20041021 | US 2004-838577 | 20040504 |
| PRAI | GB 1996-23233 | A | 19961107 | | |
| | EP 1997-950137 | A3 | 19971104 | | |
| | WO 1997-EP6180 | W | 19971104 | | |
| | US 1999-284887 | B1 | 19990527 | | |
| | US 2001-827785 | B1 | 20010406 | | |

AB The invention provides a **diphtheria**, **tetanus** and
pertussis vaccine comprising a low dose of
each of **diphtheria toxoid (D)**, **tetanus**
toxoid (T), **pertussis toxin (PT)**, filamentous
hemagglutinin (FHA) and pertactin (69K). The **vaccine** maintains
an ability to prevent **pertussis** while showing exceptionally low

reactogenicity. Combination **vaccines** comprising addnl. antigens are also provided.

RE.CNT 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

ALL CITATIONS AVAILABLE IN THE RE FORMAT

- L16 ANSWER 4 OF 43 BIOSIS COPYRIGHT (c) 2006 The Thomson Corporation on STN
AN 1999:252977 BIOSIS
DN PREV199900252977
TI Immunogenicity and reactogenicity of a **reduced-dose diphtheria-tetanus-acellular pertussis vaccine** (dTpa), a **reduced-dose acellular pertussis vaccine** (pa), and a licensed **diphtheria-tetanus (dT) vaccine** in 10-14 year-old adolescents.
AU Mertsola, J. [Reprint author]; Melot, V.; Ramalho, A.; Kaufhold, A.
CS Turku Univ. Hosp., Turku, Finland
SO Abstracts of the Interscience Conference on Antimicrobial Agents and Chemotherapy, (1998) Vol. 38, pp. 294. print.
Meeting Info.: 38th Interscience Conference on Antimicrobial Agents and Chemotherapy. San Diego, California, USA. September 24-27, 1998. American Society for Microbiology.
DT Conference; (Meeting)
Conference; Abstract; (Meeting Abstract)
Conference; (Meeting Slide)
LA English
ED Entered STN: 2 Jul 1999
Last Updated on STN: 2 Jul 199
- L16 ANSWER 5 OF 43 BIOSIS COPYRIGHT (c) 2006 The Thomson Corporation on STN
AN 1999:258918 BIOSIS
DN PREV199900258918
TI Immunogenicity and reactogenicity of a **low-content diphtheria-tetanus-acellular pertussis (DTPa) vaccine as booster** in 4 to 7-year-old children.
AU Dagan, R. [Reprint author]; Igbaria, K.; Piglansky, L.; Melot, V.; Van Brusteghem, F.; Kaufhold, A.
CS Ben-Gurion Univ., Beer-Sheva, Israel
SO Abstracts of the Interscience Conference on Antimicrobial Agents and Chemotherapy, (1998) Vol. 38, pp. 292. print.
Meeting Info.: 38th Interscience Conference on Antimicrobial Agents and Chemotherapy. San Diego, California, USA. September 24-27, 1998. American Society for Microbiology.
DT Conference; (Meeting)
Conference; Abstract; (Meeting Abstract)
Conference; (Meeting Slide)
LA English
ED Entered STN: 2 Jul 1999
Last Updated on STN: 2 Jul 1999
- L16 ANSWER 32 OF 43 BIOSIS COPYRIGHT (c) 2006 The Thomson Corporation on STN
DUPLICATE 13
AN 1985:434526 BIOSIS
DN PREV198580104518; BA80:104518
TI COMPARATIVE STUDY OF IMMUNOGLOBULINS AND SPECIFIC ANTIBODIES IN THE BLOOD OF RABBITS IMMUNIZED WITH ADSORBED **DIPHThERIA-PTERTUSSIS -TETANUS VACCINE** AN ADSORBED **DIPHThERIA-TETANUS TOXOIDS** WITH NORMAL AND **REDUCED** ANTIGEN CONTENT.
AU KOZLOVA N N [Reprint author]; GERVAZIEVA V B; IL'NITSKAYA E A; MOSHIASHVILI I YA
CS II MECHNIKOV CENT RES INST VACC SERA, MOSCOW, USSR
SO Zhurnal Mikrobiologii Epidemiologii i Immunobiologii, (1985) No. 4, pp. 39-42.
CODEN: ZMEIAV. ISSN: 0372-9311.
DT Article

FS BA
LA RUSSIAN
AB Adsorbed **DPT vaccine** and adsorbed **DT toxoids** with normal and **reduced** antigen content were used for the immunization of rabbits. The levels of IgM and IgG and the dynamics of antibodies to **diphtheria** and **tetanus** toxins and to *Bordetella pertussis* in the blood sera of the animals were studied in the postvaccinal period (on days 15 and 34). This study revealed that the **reduction** of the antigen content of adsorbed **DT toxoid** to 5 Lf of **diphtheria toxoid** and 5 binding units of **tetanus toxoid** did not decrease the capacity of the preparation for increasing the levels of IgG and IgM antibodies to **diphtheria** and **tetanus toxins** in the sera of the rabbits. The **reduced** content of these **toxoids** in adsorbed **DPT vaccine** did not affect its capacity for inducing the enhanced synthesis of IgG antibodies to **diphtheria** and **tetanus toxins**, while the production of IgM and IgA remained unchanged. At the same time an increase in the titers of antibodies to *B. pertussis* in the animals was less pronounced than that observed after the injection of commercial adsorbed **DPT vaccine**. Additional investigations are necessary in order to establish the protective potency of the **pertussis** component in adsorbed **DPT vaccine** with the **reduced** content of **toxoids** and to find out the optimum antigenic composition for this preparation.

L16 ANSWER 38 OF 43 MEDLINE on STN
AN 76129653 MEDLINE
DN PubMed ID: 1251161
TI [Booster vaccination with a **diphtheria-tetanus vaccine** for dermo-jet with low **diphtheria toxoid** content].
Auffrisch-Impfung mit einem Diphtherie-Tetanus-Impfstoff für Dermo-Jet mit niedrigem Diphtherietoxoidgehalt.
AU Wegmann A; Heiz R; Baumann T
SO Schweizerische medizinische Wochenschrift, (1976 Jan 24) Vol. 106, No. 4, pp. 112-4.
Journal code: 0404401. ISSN: 0036-7672.
CY Switzerland
DT Journal; Article; (JOURNAL ARTICLE)
LA German
FS Priority Journals
EM 197604
ED Entered STN: 19900313
Last Updated on STN: 19900313
Entered Medline: 19760427
AB Indications are provided on how the organization of polyval vaccinations against tuberculosis, **poliomyelitis** and **diphtheria-tetanus** in school collectivities can be rationalized. 19 persons aged 19-20 years were vaccinated intradermally with 1.5 Lf **diphtheria** toxoid and 15 Lf **tetanus** toxoid (contained in 0.1 ml) by means of a single injection with a jet injector (Dermo-Jet). In cases which had not previously been immunized, vaccination with the toxoid doses employed did not induce detectable antitoxin titers. In all cases which had previously been immunized the antitoxin titers after the **booster** injection were at least 400 times higher than the protection threshold for **diphtheria** and 1700 times higher than the protection threshold for **tetanus**. This was also the case in preimmunized cases which had no detectable antitoxin titer before the vaccination. The increase in antitoxin titer was inversely proportional to the prevaccinal titer in the manner of a logarithmic exponential function. It may be concluded from these results that the benefit of a **booster** vaccination is particularly high in poorly immunized persons whereas it is clearly limited in cases showing a high prevaccinal titer. The well tolerated intradermal vaccination with the jet injector (Dermo-Jet) can be considered as equivalent to the subcutaneous technic. From the point of view of organization (time consumption, problems of sterilization) this method is much preferable to intramuscular or

subcutaneous vaccination.

L16 ANSWER 41 OF 43 BIOSIS COPYRIGHT (c) 2006 The Thomson Corporation on
STN
AN 1973:219893 BIOSIS
DN PREV197356049858; BA56:49858
TI IMMUNIZATION AGAINST **DIPHTHERIA TETANUS AND**
PERTUSSIS WITH REDUCED DOSES OF ASSOCIATED
VACCINE.
AU POLIKAR A; SOLOMONOVA K; MIHAILOVA V; RUMENOVA I
SO Epidemiologiya Mikrobiologiya i Infektsiozni Bolesti, (1971)
Vol. 8, No. 1, pp. 17-21.
CODEN: EMIBA3. ISSN: 0425-1482.
DT Article
FS BA
LA Unavailable

L20 ANSWER 2 OF 15 MEDLINE on STN
 AN 95076232 MEDLINE
 DN PubMed ID: 7984981
 TI Immune status and **booster** effects of **low** doses of **tetanus toxoid** in Swedish medical personnel.
 AU Bjorkholm B; Wahl M; Granstrom M; Hagberg L
 CS Department of Infectious Diseases, Goteborg University, Ostra Hospital, Sweden.
 SO Scandinavian journal of infectious diseases, (1994) Vol. 26, No. 4, pp. 471-5.
 Journal code: 0215333. ISSN: 0036-5548.
 CY Sweden
 DT Journal; Article; (JOURNAL ARTICLE)
 LA English
 FS Priority Journals
 EM 199501
 ED Entered STN: 19950116
 Last Updated on STN: 19950116
 Entered Medline: 19950105
 AB Of 102 medical staff at a Swedish hospital, 81% had tetanus antitoxin titres > or = 0.01 IU/ml in 1984-85. The unprotected individuals (antitoxin titre < 0.01 IU/ml) were all > 30 years of age. Of this group, one-third lacked a protective antibody level against tetanus toxin. **Low booster** doses of **tetanus toxoid** (0.75 or 1.9 Lf) were given to 66 vaccinees with a history of previous basic vaccination and no history of **booster** vaccination within the previous 5 years. The median titre increased from 0.26 IU/ml before to 3.3 IU/ml after vaccination. **Low** doses of **tetanus toxoid** may thus still provide an adequate immune response when given as a **booster** vaccination to individuals with a reliable history of basic immunization.

L20 ANSWER 3 OF 15 MEDLINE on STN
 AN 90069457 MEDLINE
 DN PubMed ID: 2587945
 TI Immune status and **booster** effects of **low** doses of **diphtheria toxoid** in Swedish medical personnel.
 AU Bjorkholm B; Wahl M; Granstrom M; Hagberg L
 CS Department of Infectious Diseases, University of Goteborg, Ostra Hospital, Sweden.
 SO Scandinavian journal of infectious diseases, (1989) Vol. 21, No. 4, pp. 429-34.
 Journal code: 0215333. ISSN: 0036-5548.
 CY Sweden
 DT Journal; Article; (JOURNAL ARTICLE)
 LA English
 FS Priority Journals
 EM 198912
 ED Entered STN: 19900328
 Last Updated on STN: 19900328
 Entered Medline: 19891222
 AB During a diphtheria outbreak among Swedish alcoholics in 1984-1985, only 60% of 328 medical staff at risk for exposure had diphtheria antitoxin titers greater than or equal to 0.01 IU/ml, which is usually considered to give relative protection. 21% had levels between 0.01-0.09 IU/ml and the remaining 39% had titers greater than or equal to 0.1 IU/ml. **Booster** doses of 0.5 Lf-12.5 Lf of diphtheria toxoid were given to 450 vaccinees. Of 72 individuals with **low** pre-immunization titers, who were immunized with less than or equal to 2.5 Lf **diphtheria toxoid**, 40% failed to reach greater than or equal to 0.1 IU/ml when analyzed 4 weeks after vaccination. Local tenderness and swelling greater than 5 cm at the site of injection or general discomfort was found in 11% of those immunized with **low** dose **diphtheria toxoid** (less than or equal to 2.5 Lf). When the dose 0.5 Lf of diphtheria toxoid was combined with tetanus toxoid (3.75 Lf) the frequency of adverse reactions increased to 34% (p less than 0.001). The study shows that vaccination status in medical personnel must also be continuously examined in countries where **diphtheria** is

'rare and that **low booster** doses of **diphtheria toxoid** may not achieve an adequate immune response.

L20 ANSWER 5 OF 15 MEDLINE on STN
AN 81172425 MEDLINE
DN PubMed ID: 7219283
TI **Diphtheria** immunization in adolescents and adults with **reduced** doses of adsorbed **diphtheria toxoid**.
AU Feery B J; Benenson A S; Forsyth J R; Menser M A; Minty D W
SO The Medical journal of Australia, (1981 Feb 7) Vol. 1, No. 3, pp. 128-30.
Journal code: 0400714. ISSN: 0025-729X.
CY Australia
DT Journal; Article; (JOURNAL ARTICLE)
LA English
FS Priority Journals
EM 198106
ED Entered STN: 19900316
Last Updated on STN: 19900316
Entered Medline: 19810613
AB A study was undertaken in Schick-positive university students and schoolchildren to determine the **diphtheria** antitoxin response to **vaccines** containing a **reduced** dose of adsorbed **diphtheria toxoid**. It was found that the majority of participants, apparently previously sensitized, responded with an increase in antitoxin titre to protective levels after performance of the Schick test, or after the first dose of **vaccine**. A group of non-immune students required three doses of **vaccine** to reach adequate antitoxin levels to ensure durable immunity. Only one local reaction was observed in the group of 51 students, and this was attributed to an Arthus-type reaction involving the tetanus toxoid component of a combined adsorbed diphtheria and tetanus **vaccine**.

L20 ANSWER 8 OF 15 MEDLINE on STN
AN 69090769 MEDLINE
DN PubMed ID: 5705656
TI [The usefulness of **boosters** of nonadsorbed **diphtheria toxoid** in **reduced** doses].
Opportunité des rappels d'anatoxine diphtérique non adsorbée et à doses réduites.
AU Joubert L; Fauchet S; Picard F; Triau R
SO Revue d'immunologie et de thérapie antimicrobienne, (1968 Jul-Sep) Vol. 32, No. 4, pp. 245-51.
Journal code: 0253026. ISSN: 0035-2454.
CY France
DT Journal; Article; (JOURNAL ARTICLE)
LA French
FS Priority Journals
EM 196903
ED Entered STN: 19900101
Last Updated on STN: 19950206
Entered Medline: 19690306

L20 ANSWER 12 OF 15 BIOSIS COPYRIGHT (c) 2006 The Thomson Corporation on STN
AN 1995:275303 BIOSIS
DN PREV199598289603
TI Can **reductions** in **diphtheria toxoid** or **aluminum** content **reduce** the reactogenicity of **booster** doses of **DPT vaccine**?
AU Scheifele, David W. [Reprint author]; Meekison, William; Guasparini, Roland; Barreto, Luis
CS Vaccine Eval. Cent., 950 West 28th Ave., Vancouver, BC V5Z 4H4, Canada
SO Immunology and Infectious Diseases (Oxford), (1995) Vol. 5, No. 1, pp. 73-77.
CODEN: IINDEK. ISSN: 0959-4957.
DT Article

LA English
ED Entered STN: 26 Jun 1995
Last Updated on STN: 26 Jun 1995
AB Large local reactions often follow **booster** doses of the diphtheria-pertussis-tetanus (DPT) **vaccine** used in Canada. Their cause is uncertain but the relatively large dose of diphtheria toxoid is a possible contributor. This study was designed to determine if **vaccines** containing less diphtheria toxoid (25, 15 or 10 Lf per dose) and/or aluminum (0.33 or 0.17 mg per dose) would cause fewer adverse effects. Subjects included 373 healthy 4- to 6-year-olds due for a fifth dose of DPT **vaccine**. They were randomly assigned to receive one of five DPT **vaccines**, in double-blinded fashion. Local and systemic adverse reactions were assessed 24 h post-immunization by visiting research nurses. Blood was tested for antitoxins prior to and 4 weeks after immunization. The lot of standard **vaccine** caused fever in 21.9% and gtoreq 50 mm of injection site redness in 49.3% and swelling in 41.0%. Limited arm movement was detected in 26% and was severe in 5.5%. None of the modified **vaccines** reduced such morbidity significantly. All formulations produced large increases in tetanus and diphtheria antitoxin levels. It is concluded that the principal cause of local reactions after **vaccine** is not diphtheria toxoid or aluminum but a component(s) that was not varied among the five preparations, such as pertussis **vaccine**. The advent of acellular pertussis **vaccines** might provide a remedy.

L20 ANSWER 13 OF 15 BIOSIS COPYRIGHT (c) 2006 The Thomson Corporation on STN
AN 1981:295875 BIOSIS
DN PREV198172080859; BA72:80859
TI DIPHTHERIA IMMUNIZATION IN ADOLESCENTS AND ADULTS WITH REDUCED DOSES OF ADSORBED DIPHTHERIA TOXOID.
AU FEERY B J [Reprint author]; BENENSON A S; FORSYTH J R L; MENSER M A; MINTY D W
CS COMMONW SERUM LAB, 45 POPLAR RD, PARKVILLE, VIC 3052
SO Medical Journal of Australia, (1981) Vol. 68-1, No. 3, pp. 128-130.
CODEN: MJAUAJ. ISSN: 0025-729X.
DT Article
FS BA
LA ENGLISH
AB A study was undertaken in Schick-positive university students and schoolchildren to determine the **diphtheria** antitoxin response to **vaccines** containing a **reduced** dose of adsorbed **diphtheria toxoid**. The majority of participants, apparently previously sensitized, responded with an increase in antitoxin titer to protective levels after performance of the Schick test or after the 1st dose of **vaccine**. A group of non-immune students required 3 doses of **vaccine** to reach adequate antitoxin levels to ensure durable immunity. Only 1 local reaction was observed in the group of 51 students and this was attributed to an Arthus-type reaction involving the tetanus toxoid component of a combined adsorbed diphtheria and tetanus **vaccine**.